

10/629,975
Search
LYCOK 6/27/07

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(FILE 'HOME' ENTERED AT 11:55:06 ON 27 JUN 2007)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 11:55:24 ON 27
JUN 2007

| | |
|----|--|
| L1 | 89 S (IBD TREATMENT) |
| L2 | 7 S L1 AND PD<2001 |
| L3 | 5 DUPLICATE REMOVE L2 (2 DUPLICATES REMOVED) |
| L4 | 0 S L3 AND LACTOFER? |
| L5 | 0 S L3 AND LEUKOC? |
| L6 | 4 S (MONITOR? IBD) |
| L7 | 2 DUPLICATE REMOVE L6 (2 DUPLICATES REMOVED) |

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| L7 | 2 DUPLICATE REMOVE L6 (2 DUPLICATES REMOVED) |

=>

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

AN 1995:522917 CAPLUS
 DN 122:260562
 ED Entered STN: 04 May 1995
 TI Diagnostic test and kit for disease or disorders in the digestive system
 IN Fagerhol, Magne K.; Dale, Inge; Roseth, Arne G.
 PA Norway
 SO Can. Pat. Appl.
 CODEN: CPXXEB
 DT Patent
 LA English
 IC ICM G01N033-68
 ICS G01N033-574
 CC 9-10 (Biochemical Methods)
 Section cross-reference(s): 14, 15

FAN.CNT 1

| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------|---------------|------|----------|-----------------|----------|
| PI | CA 2123856 | A1 | 19941128 | CA 1994-2123856 | 19940518 |
| | US 5455160 | A | 19951003 | US 1993-67802 | 19930527 |
| PRAI | US 1993-67802 | A | 19930527 | | |

CLASS

| PATENT NO. | CLASS | PATENT FAMILY CLASSIFICATION CODES |
|------------|-------|---|
| CA 2123856 | ICM | G01N033-68 |
| | ICS | G01N033-574 |
| | IPCI | G01N0033-68 [ICM,5]; G01N0033-574 [ICS,5] |
| | IPCR | G01N0033-574 [I,C*]; G01N0033-574 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A] |
| US 5455160 | IPCI | G01N0033-53 [ICM,6]; G01N0033-574 [ICS,6] |
| | IPCR | G01N0033-574 [I,C*]; G01N0033-574 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A] |
| | NCL | 435/007.230; 435/007.920; 435/007.930; 435/961.000; 436/064.000; 436/811.000; 436/813.000 |
| | ECLA | G01N033/574K; G01N033/68 |

AB This study describes a method for extraction and quantification of calprotectin (L1 protein) in feces by enzyme immunoassay. This protein is a prominent antimicrobial component of neutrophils, monocytes, macrophages and squamous epithelia. Calprotectin was stable in feces during storage for 7 days at room temperature. Fecal calprotectin quantitated in a 5-g sample taken from a 24-h feces collection gave a reliable estimate of calprotectin found in the pooled collection. The assay had a within assay precision (CV) of 1.9% and a between assay precision of 14.8%. The following mean fecal calprotectin levels were found: healthy subjects 3095 µg/L; hospital controls 14,637 µg/L; and patients with inflammatory bowel disease (Crohn's disease and ulcerative colitis) 40,850 µg/L. The difference between the means are highly significant. All patients with IBD and 10 of 11 patients with gastrointestinal carcinomas had calprotectin level above the suggested reference limit of 9000 µg/L. Determination of fecal

calprotectin is

an important routine parameter for monitoring IBD and gastrointestinal cancer.

ST feces calprotectin detn diagnosis disease cancer; enzyme immunoassay calprotectin feces; inflammatory bowel disease diagnosis calprotectin; gastrointestinal tract calprotectin detn; colorectal carcinoma diagnosis calprotectin detn

IT Feces

(calprotectin determination in feces by EIA in diagnosis of disease or disorders of digestive system)

IT Antibodies

RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(calprotectin determination in feces by EIA in diagnosis of disease or

disorders of digestive system)

IT Bone marrow
 (depression; calprotectin determination in feces by EIA in diagnosis of
 disease
 or disorders of digestive system)

IT Intestine, disease
 (Crohn's, calprotectin determination in feces by EIA in diagnosis of
 disease or
 disorders of digestive system)

IT Proteins, specific or class
 RL: ANT (Analyte); THU (Therapeutic use); ANST (Analytical study); BIOL
 (Biological study); USES (Uses)
 (L1, calprotectin determination in feces by EIA in diagnosis of disease or
 disorders of digestive system)

IT Digestive tract
 (disease, calprotectin determination in feces by EIA in diagnosis of
 disease or
 disorders of digestive system)

IT Intestine, disease
 (inflammatory, calprotectin determination in feces by EIA in diagnosis of
 disease or disorders of digestive system)

IT Intestine, neoplasm
 (large, carcinoma, calprotectin determination in feces by EIA in diagnosis
 of
 disease or disorders of digestive system)

IT Digestive tract
 (neoplasm, carcinoma, calprotectin determination in feces by EIA in
 diagnosis
 of disease or disorders of digestive system)

IT Intestine, disease
 (ulcerative colitis, calprotectin determination in feces by EIA in
 diagnosis of
 disease or disorders of digestive system)

=>

ANSWER 1 OF 5 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN
 DUPLICATE 1

AN 2001:33185 BIOSIS
 DN PREV200100033185
 TI Natalizumab. Treatment of IBD, treatment of multiple
 sclerosis: AN100226, AntegrenTM.
 AU Sorbera, L. A. [Reprint author]; Martin, L. [Reprint author]; Rabasseda,
 X. [Reprint author]
 CS Prous Science, 08080, Barcelona, Spain
 SO Drugs of the Future, (September, 2000) Vol. 25, No. 9, pp.
 917-921. print.
 ISSN: 0377-8282.

DT Article
 LA English
 ED Entered STN: 10 Jan 2001
 Last Updated on STN: 12 Feb 2002

CC Pharmacology - Clinical pharmacology 22005
 Pathology - Therapy 12512
 Digestive system - Pathology 14006
 Nervous system - Pathology 20506
 Pharmacology - General 22002
 Pharmacology - Immunological processes and allergy 22018
 Immunology - General and methods 34502
 Immunology - Immunopathology, tissue immunology 34508

IT Major Concepts
 Clinical Immunology (Human Medicine, Medical Sciences);
 Gastroenterology (Human Medicine, Medical Sciences); Pharmacology

IT Diseases
 inflammatory bowel disease: digestive system disease
 Inflammatory Bowel Diseases (MeSH)

IT Diseases
 multiple sclerosis: immune system disease, nervous system disease
 Multiple Sclerosis (MeSH)

IT Chemicals & Biochemicals
 natalizumab [AN 100226, Antegren]: immunosuppressant-drug, monoclonal
 antibody

ORGN Classifier
 Hominidae 86215
 Super Taxa
 Primates; Mammalia; Vertebrata; Chordata; Animalia
 Organism Name
 human
 Taxa Notes
 Animals, Chordates, Humans, Mammals, Primates, Vertebrates

RN 189261-10-7 (natalizumab)
 189261-10-7 (AN 100226)
 189261-10-7 (Antegren)

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FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 17:26:06 ON 27 JUN 2007

L1 66455 S (ULCERATIVE COLITIS)
L2 248 S L1 AND LACTOFERRIN?
L3 23 S L2 AND TREATMENT?
L4 17 DUPLICATE REMOVE L3 (6 DUPLICATES REMOVED)
L5 7 S L4 AND PD<2001
L6 35 S L2 AND THERAPY
L7 35 S L6 NOT L5
L8 27 DUPLICATE REMOVE L7 (8 DUPLICATES REMOVED)
L9 9 S L8 AND PD<2001

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L7 35 S L6 NOT L5
L8 27 DUPLICATE REMOVE L7 (8 DUPLICATES REMOVED)
L9 9 S L8 AND PD<2001

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reserved on STN

AN 1999322801 EMBASE

TI Faecal parameters in the assessment of activity in inflammatory bowel disease.

AU Van der Sluys Veer A.; Biemond I.; Verspaget H.W.; Lamers C.B.H.W.

CS A. Van der Sluys Veer, Dept of Gastroenterol. and Hepatol., Leiden University Medical Center, Building 1, PO Box 9600, C4-PNL-2300 RC Leiden, Netherlands

SO Scandinavian Journal of Gastroenterology, Supplement, (1999)
Vol. 33, No. 230, pp. 106-110.

Refs: 55

ISSN: 0085-5928 CODEN: SJGSB8

CY Norway

DT Journal; Article

FS 005 General Pathology and Pathological Anatomy

029 Clinical Biochemistry

037 Drug Literature Index

048 Gastroenterology

LA English

SL English

ED Entered STN: 30 Sep 1999

Last Updated on STN: 30 Sep 1999

AB Background: Determination of inflammatory activity is helpful when assessing the efficacy of drugs in therapeutic trials and in facilitating management of individual patients with inflammatory bowel disease (IBD). Faecal parameters have been hypothesized to be more specific than non-faecal measurements in the assessment of intestinal inflammation. Methods: Review of the literature on faecal measurements in IBD. Results and conclusions: Leakage of various proteins and leukocyte products into the intestinal lumen can be assessed and quantified in stool specimens and serve as a measurement of inflammatory activity. Several of these faecal parameters are raised in patients with IBD. There is a considerable overlap between patients with active and those with inactive disease, however, and the correlation of the faecal parameters with disease activity indices is often low. The value of α -apprx.1-antitrypsin measurement in faeces in the assessment of intestinal inflammation has been well established. Further studies in patients with IBD are needed to determine whether other faecal parameters, such as lactoferrin, tumour necrosis factor α , PMN-elastase, lysozyme, leucocyte esterase, immunoglobulin A, among others, are more accurate or cost-effective than measurement of α -apprx.1-antitrypsin in the stools of such patients.

CT Medical Descriptors:

*feces

*enteritis: DI, diagnosis

*enteritis: DT, drug therapy

*Crohn disease: DI, diagnosis

*Crohn disease: DT, drug therapy

*ulcerative colitis: DI, diagnosis

*ulcerative colitis: DT, drug therapy

gastrointestinal endoscopy

intestine biopsy

imaging

immunodiffusion

enzyme immunoassay

nephelometry

human

clinical trial

article

priority journal

Drug Descriptors:

*alpha 1 antitrypsin: EC, endogenous compound

protein: EC, endogenous compound

ANSWER 6 OF 9 EMBASE COPYRIGHT (c) 2007 Elsevier B.V. All rights

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*Crohn disease: DI, diagnosis

*Crohn disease: DT, drug therapy

*ulcerative colitis: DI, diagnosis

*ulcerative colitis: DT, drug therapy

gastrointestinal endoscopy

intestine biopsy

imaging

immunodiffusion

enzyme immunoassay

nephelometry

human

clinical trial

article

priority journal

Drug Descriptors:

*alpha 1 antitrypsin: EC, endogenous compound

protein: EC, endogenous compound

lactoferrin: EC, endogenous compound
tumor necrosis factor alpha: EC, endogenous compound
leukocyte elastase: EC, endogenous compound
lysozyme: EC, endogenous compound
esterase: EC, endogenous compound
immunoglobulin a: EC, endogenous compound
barium
methylprednisolone: CB, drug combination
methylprednisolone: DT, drug therapy
salazosulfapyridine: CB, drug combination
salazosulfapyridine: DT, drug therapy
hemoglobin: EC, endogenous compound
indium 111

RN (alpha 1 antitrypsin) 9041-92-3; (protein) 67254-75-5; (
lactoferrin) 55599-62-7; (leukocyte elastase) 109968-22-1;
(lysozyme) 9001-63-2; (esterase) 9013-79-0; (barium) 7440-39-3;
(methylprednisolone) 6923-42-8, 83-43-2; (salazosulfapyridine) 599-79-1;
(hemoglobin) 9008-02-0; (indium 111) 15750-15-9

lactoferrin: EC, endogenous compound
tumor necrosis factor alpha: EC, endogenous compound
leukocyte elastase: EC, endogenous compound
lysozyme: EC, endogenous compound
esterase: EC, endogenous compound
immunoglobulin a: EC, endogenous compound
barium
methylprednisolone: CB, drug combination
methylprednisolone: DT, drug therapy
salazosulfapyridine: CB, drug combination
salazosulfapyridine: DT, drug therapy
hemoglobin: EC, endogenous compound
indium 111

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(lysozyme) 9001-63-2; (esterase) 9013-79-0; (barium) 7440-39-3;
(methylprednisolone) 6923-42-8, 83-43-2; (salazosulfapyridine) 599-79-1;
(hemoglobin) 9008-02-0; (indium 111) 15750-15-9

ANSWER 4 OF 9 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN

AN 1993:396686 BIOSIS

DN PREV199345055511

TI A study to determine fecal lactoferrin in patients with
ulcerative colitis.

AU Sudo, Ichiro

CS Fourth Dep. Intern. Med., Tokyo Med. Coll., Japan

SO Japanese Journal of Gastroenterology, (1993) Vol. 90, No. 4, pp.
824.
ISSN: 0446-6586.

DT Article

LA Japanese

ED Entered STN: 30 Aug 1993
Last Updated on STN: 30 Aug 1993

CC Biochemistry studies - Proteins, peptides and amino acids 10064
Pathology - Diagnostic 12504
Pathology - Inflammation and inflammatory disease 12508
Pathology - Therapy 12512
Digestive system - General and methods 14001
Digestive system - Pathology 14006
Immunology - Immunopathology, tissue immunology 34508

IT Major Concepts
Clinical Endocrinology (Human Medicine, Medical Sciences); Digestive
System (Ingestion and Assimilation); Gastroenterology (Human Medicine,
Medical Sciences); Pathology

IT Miscellaneous Descriptors
DIAGNOSTIC METHOD; PATHOLOGY; THERAPY

ORGN Classifier
Hominidae 86215
Super Taxa
Primates; Mammalia; Vertebrata; Chordata; Animalia
Organism Name
human
Taxa Notes
Animals, Chordates, Humans, Mammals, Primates, Vertebrates

ANSWER 4 OF 9 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN

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SO Japanese Journal of Gastroenterology, (1993) Vol. 90, No. 4, pp.
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DT Article

LA Japanese

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Last Updated on STN: 30 Aug 1993

CC Biochemistry studies - Proteins, peptides and amino acids 10064
Pathology - Diagnostic 12504
Pathology - Inflammation and inflammatory disease 12508
Pathology - Therapy 12512
Digestive system - General and methods 14001
Digestive system - Pathology 14006
Immunology - Immunopathology, tissue immunology 34508

IT Major Concepts
Clinical Endocrinology (Human Medicine, Medical Sciences); Digestive
System (Ingestion and Assimilation); Gastroenterology (Human Medicine,
Medical Sciences); Pathology

IT Miscellaneous Descriptors
DIAGNOSTIC METHOD; PATHOLOGY; THERAPY

ORGN Classifier
Hominidae 86215
Super Taxa
Primates; Mammalia; Vertebrata; Chordata; Animalia
Organism Name
human
Taxa Notes
Animals, Chordates, Humans, Mammals, Primates, Vertebrates

ANSWER 1 OF 9 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN

AN 1994:286260 BIOSIS

DN PREV199497299260

TI Mucosal lactoferrin, "sticky" neutrophils, and pathergy in
ulcerative colitis and pyoderma gangrenosum:
Implications for pathogenesis, and successful therapy with
heparin.

AU Dwarakanath, A. D. [Reprint author]; Finnie, I. A. [Reprint author]; Yu,
L. G. [Reprint author]; O'Dowd, G. M.; Rhodes, Jonathan M.

CS Dep. Med., Univ. Liverpool, P.O. Box 147, Liverpool L69 3BX, UK

SO Gastroenterology, (1994) Vol. 106, No. 4 SUPPL., pp. A674.
Meeting Info.: 95th Annual Meeting of the American Gastroenterological
Association. New Orleans, Louisiana, USA. May 15-18, 1994.
CODEN: GASTAB. ISSN: 0016-5085.

DT Conference; (Meeting)
Conference; Abstract; (Meeting Abstract)

LA English

ED Entered STN: 30 Jun 1994
Last Updated on STN: 1 Jul 1994

CC General biology - Symposia, transactions and proceedings 00520
Cytology - Human 02508
Biochemistry studies - Carbohydrates 10068
Biophysics - Molecular properties and macromolecules 10506
Pathology - Inflammation and inflammatory disease 12508
Pathology - Therapy 12512
Digestive system - Pathology 14006
Blood - Blood cell studies 15004
Blood - Blood, lymphatic and reticuloendothelial pathologies 15006
Blood - Lymphatic tissue and reticuloendothelial system 15008
Pharmacology - Clinical pharmacology 22005
Pharmacology - Digestive system 22014

IT Major Concepts
Biochemistry and Molecular Biophysics; Blood and Lymphatics (Transport
and Circulation); Cell Biology; Gastroenterology (Human Medicine,
Medical Sciences); Hematology (Human Medicine, Medical Sciences);
Pathology; Pharmacology

IT Chemicals & Biochemicals
HEPARIN

IT Miscellaneous Descriptors
GASTROINTESTINAL-DRUG; HEPARIN; MEETING ABSTRACT

ORGN Classifier
Hominidae 86215
Super Taxa
Primates; Mammalia; Vertebrata; Chordata; Animalia
Organism Name
human
Taxa Notes
Animals, Chordates, Humans, Mammals, Primates, Vertebrates

RN 9005-49-6 (HEPARIN)



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Matcher

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NLM Gateway

TOXNET

Consumer Health

Clinical Alerts

ClinicalTrials.gov

PubMed Central

1: Clin Biochem. 1994 Aug;27(4):259-64.

ELSEVIER
FULL-TEXT ARTICLE

Related Articles, Links

Immunochemical detection of human lactoferrin in feces as a new marker for inflammatory gastrointestinal disorders and colon cancer.

Uchida K, Matsuse R, Tomita S, Sugi K, Saitoh O, Ohshiba S.

Kyoto Medical Science Laboratory, Kyoto, Japan.

We have developed a new immunochemical test for fecal lactoferrin (LF) utilizing an enzyme-linked immunosorbent assay (ELISA). The ELISA had a sensitivity of about 10 micrograms/L of lactoferrin and the measurable range was 10.0-1000.0 micrograms/L (1.0-100.0 micrograms LF/g feces). The stability of lactoferrin in feces was greater than that of myeloperoxidase and leucocyte elastase. The fecal concentration of lactoferrin (mean +/- SD) in 35 normal subjects was 0.75 +/- 0.83 microgram/g feces, whereas that in 24 patients with colon cancer was 74.4 +/- 88.3 micrograms/g feces. The fecal lactoferrin concentration of 38 patient with active ulcerative colitis was 307.4 +/- 233.9 micrograms/g feces, and that in 36 patients with active Crohn's disease was 191.7 +/- 231.1 micrograms/g feces. The ELISA for human fecal lactoferrin might be useful in the diagnosis of colon disease.

Publication Types:

- Comparative Study

MeSH Terms:

- Adolescent
- Adult
- Aged
- Colitis, Ulcerative/diagnosis
- Colitis, Ulcerative/metabolism*
- Colonic Neoplasms/diagnosis
- Colonic Neoplasms/metabolism*
- Colonic Polyps/diagnosis
- Colonic Polyps/metabolism
- Colonoscopy
- Crohn Disease/diagnosis
- Crohn Disease/metabolism*
- Enzyme-Linked Immunosorbent Assay
- Feces/chemistry*
- Female
- Humans
- Lactoferrin/blood

- [Lactoferrin/metabolism*](#)
- [Male](#)
- [Middle Aged](#)
- [Regression Analysis](#)

Substances:

- [Lactoferrin](#)

PMID: 8001286 [PubMed - indexed for MEDLINE]

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Department of Health & Human Services

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